Appl. No. 10/552,134

Amdt. Dated: October 28, 2010

Reply to Office Action of June 28, 2010

Amendments to the Claims.

This listing of claims will replace all prior versions, and listings, of claims in the application.

Listing of Claims:

- 1. (Currently amended) A method of producing a radiolabelled gallium complex in a form
 - suitable for use in PET or SPECT radiopharmaceutical imaging, said method comprising

by reacting a Ga³⁺ radioisotope in a suitable solvent with a macrocyclic bifunctional

chelating agent, wherein said macrocyclic bifunctional chelating agent is linked to a

targeting vector selected from the group consisting of proteins, glycoproteins,

lipoproteins, polypeptides, glycopolypeptides, lipopolypeptides, peptides, glycopeptides,

lipopeptides, carbohydrates, nucleic acids, oligonucleotides or small organic molecules;

characterised in that the reaction is carried out using microwave activation at 80 to

120 W for 20 s to 2 min.

- 2. (Previously presented) The method according to claim 1 wherein the Ga³⁺ radioisotope is selected from the group consisting of 66Ga3+, 67Ga3+ and 68Ga3+.
- 3. (Previously presented) The method according to claim 1 wherein the Ga³⁺ radioisotope is 68Ga3+
- 4. (Cancelled)
- 5. (Previously presented) The method according to claim 1 wherein the macrocyclic bifunctional chelating agent comprises hard donor atoms, preferably O and N atoms.
- (Cancelled)

Appl. No. 10/552,134

Amdt. Dated: October 28, 2010

Reply to Office Action of June 28, 2010

7. (Cancelled)

8. (Previously presented) The method according to claim 1 wherein the target vector is a

peptide or oligonucleotide.

9. (Previously presented) The method according to claim 1 wherein the microwave

activation is carried out at 90 to 110 W.

10. (Previously presented) The method according to claim 1 wherein the microwave

activation is carried out for 30 s to 90 s.

11. (Previously presented) The method according to claim 3 wherein the ⁶⁸Ga³⁺ is obtained

by contacting the eluate from a 68Ge/68Ga generator with an anion exchanger and eluting

⁶⁸Ga³⁺ from said anion exchanger.

12. (Previously presented) The method according to claim 11 wherein the 68Ge/68Ga

generator comprises a column comprising titanium dioxide.

13. (Previously presented) The method according to claim 11 wherein the anion exchanger

comprises HCO3 as counterions.

14. (Previously presented) The method according to claim 11 wherein the anion exchanger is

an anion exchanger comprising quaternary amine functional groups, or the ion exchanger

is a anion exchange resin based on polystyrene-divinylbenzene.

15. (Previously presented) The method according to claim 1 for the production of ⁶⁸Ga-

radiolabelled PET tracers.

Page 3 of 10

Appl. No. 10/552,134

Amdt. Dated: October 28, 2010

Reply to Office Action of June 28, 2010

16. (Withdrawn) Method according to claim 11 wherein the eluting ⁶⁸Ga³⁺ is in the picomolar to nanomolar range after the elution, and more preferably in a nanomolar to micromolar level.